IN THE UNITED STATES PATENT AND TRADEMARK OFFICE Examiner Lyle A. Alexander

ART UNIT 1743

In re application of E. Alan Bates et al. Application No. 08/935,629 Filed 09/23/97 For ASSAYING DEVICE ....

### **DECLARATION UNDER 37 CFR 1.132**

Commissioner for Patents Washington, DC 20231

- I, Gary Hoffman, declare as follows:
- 1. I am the third joint inventor for this application.
- 2. While the claims of this application have been rejected based on the patent of Senior, the disclosure of the application starts from a technology actually quite different from that of Senior. Characteristic of this difference is Senior's provision of its bibulous member 16 protruding out of the housing. When the bibulous member is placed in a urine stream, it is quickly soaked. Senior has correctly chosen to display this version in its drawings, because versions such as suggested in paragraph a. in col. 5 of Senior would tend to shield the bibulous member from a urine stream.
- 3. This application references in its BACKGROUND a quite different technology, that of drug testing using an immunoassay method called antigen-antibody competitive binding. Characteristic of this technology is the measured application of only a few drops of urine. This dropwise application of the urine is disclosed in the specification in the first paragraph of the DETAILED DESCRIPTION OF THE INVENTION.
  - 4. Attached hereto are Exhibits 1 to 5 demonstrating five different instances of the type

From: Sullivan To: Exr. Lyle Alexander Date: 5/26/04 Time: 2:45:00 PM Page 9 of 33

of technology forming the starting point for this application. These are as follows:

Exhibit 1 - Pages 1-4 of a document entitled "AccuSign DOA 4, THC/OPI/COC/AMP" bearing copyright notice dated 1996;

- Exhibit 2 Pages 1-4 of a document entitled "AccuSign BAR", also bearing copyright notice dated 1996;
- Exhibit 3 Front and back of a leaflet headed "Drug Test Resources International", likewise concerning AccuSign DOA 4 and bearing a 1996 copyright notice;
- Exhibit 4 One-sided, undated leaflet headed "Visaline II"; and
- Exhibit 5 Copies of the packages of several kits labeled HOME DRUG TEST and their instruction leaflets.
- 5. Exhibit 1 is noteworthy for the correspondence of the terminology in its section Principle on its page 1 with the terminology in the BACKGROUND section of this application.
- 6. All of the exhibits direct that 3 drops of urine be applied, this being a standard for this technology. All of the exhibits provide either a dropper or a pipette for transfer of the urine sample into the sample well. In each of the exhibits, the sample receiving area is clearly a well, and Exhibit 1, for instance, calls it a "well" in the section Test Protocol on its page 2.
- 7. The chart at the top of page 2 of Exhibit 1, for instance, explains that appearance of a line for a particular drug in this starting technology is a negative indication, i.e. the drug is not present in amounts above the cutoff level.
- 8. A characteristic of this technology is that the sample collection location must not be flooded with urine, as in Senior, because this leads to false positives by washing out the lines that would otherwise indicate negative readings.

FROM : HOFFMAN

FAX NO. : 412-821-2420 Aug. 21 2000 02:40PM P1

- 9. It is to be noted that none of these exhibits mentions photocopying the results. The statements regarding photocopying in the BACKGROUND section of the present application represent perceptions of the present inventor group, rather than the state of the art at the time this invention was made.
- 10. While the provision of a cap to cover Senior's wet, protruding member 16 is immediately understandable, it was the present group of inventors which perceived the advantages of such for the different technology from which the present invention arose.
- 11. A difference between the present invention and Senior concerns the problem with putting a cap on Senior's test. Because the bibulous material can be so flexible (especially when wet) there often is difficulty slipping the cap on without bending the material. It's like threading a needle by holding the thread still and moving the needle.
- 12. All statements made herein of my own knowledge are true and all statements made on information and belief are believed to be true; such statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and may jeopardize the validity of the application or any patent issued thereon.

Signature: Qay Date: 8-21-00

### AccuSign DOA 4

### One-Step Panel Assay for Drugs of Abuse

For In Vitro Use Only

Simple One-Step Immunoassay for the Qualitative Detection of THC metabolites, Opiates, Cocaine metabolite, Amphetamines, and/or their Metabolites in Urine.

### PBM

Catalog No.

DOA-240 DOA-240-10

35 Test Kit 10 Test Kit

### Intended Use

The AccuSign" DOA 4 THC/OPI/COC/AMP Panel Assay is a simple, one-step immunochromatographic test for the rapid, qualitative detection of THC metabolites, opiates, cocaine metabolite, and amphetamines in urine. The test detects the major metabolites of these drugs at the following cutoff concentrations.

THC	11-nor-Δ*-THC-9-carboxylic acid	50 ng/mL
OPI	Morphine	300 ng/mL
COC	Benzoylecgonine	300 ng/mL
AMP	Amphetamine	1000 ng/mI.

The Accusign DOA 4 THC/OPI/COC/AMP test provides only a preliminary analytical result. A more specific alternative chemical method must be used in order to obtain a confirmed analytical result. Gas chromatography, mass spectrometry (GC/MS), is the preferred confirmatory method. Other chemical confirmatory methods are available. Clinical consideration and professional judgment should be applied to any drug of abuse test result, particularly when preliminary positive results are used.

### Summary and Explanation

Drug abuse has become one of the most destructive social problems in recent years, affecting nearly every corner of the world. To effectively combat this increasingly disturbing problem, there is a strong need for a simple, rapid, inexpensive, disposable, visual, and non-instrument requiring drug screening test kit. According to the National Institute on Drug Abuse (NIDA), THC (Marijuana), Opiates, Cocaine, and Amphetamines are among the most widely abused drugs. The one-step AccuSign\* DOA 4 Panel Assay is a test for screening these four major drugs of abuse in urine, simultaneously with one sample application. The test takes less than 10 minutes to perform.

THC ( $\Delta^{\bullet}$ -tetrahydrocannabinol) is the primary active ingredient in cannabinoids (marijuana). When ingested or smoked, it produces euphoric effects. Users experience impatrment of short term memory and THC use slows learning. Also, it may cause transient episodes of confusion, anxiety, or frank toxic delirium. Long term, relatively heavy use may be associated with behavioral disorders. The peak

effect of smoking THC occurs in 20–30 minutes and the duration is 90–120 minutes after one eigarette. Elevated levels of urinary metabolites are found within hours of exposure and remain detectable for 3–10 days after smoking. The main metabolite excreted in the urne is 11-nor- $\Delta$ -tetrahydrocannabinol-9-carboxylic acid.

Opioid analgesics comprise a large group of substances which control pain by depressing the central nervous system. Morphine is the prototype compound of this group. Morphine is excreted unmetabolized, and is also the major metabolic product of codeine and heroin. Morphine is detectable in the urine for several days after an opiate dose.

Cocaine, derived from the leaves of coca plant, is a potent central nervous system (CNS) stimulant and a local anesthetic. Cocaine induces euphoria, confidence and a sense of increased energy in the user, these psychological effects are accompanied by increased heart rate, dilation of the pupils, fever, tremors and sweating. Cocaine is used by smoking, intravenous, intranasal or oral administration, and excreted in the urne primarily as benzoylecgonine in a short time. Benzoylecgonine has a longer biological half-life (5–8 hours) than cocaine (0.5–1.5 hours) and can generally be detected for 24–60 hours after cocaine use or exposure.<sup>34</sup>

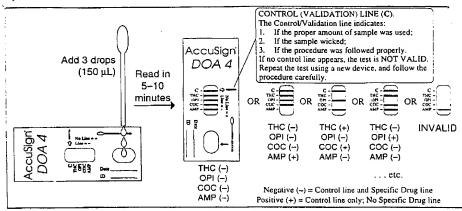
Amphetamine is a potent sympathomimetic agent with therapeutic applications. The drug can be taken orally, injected, or inhaled. Acute higher doses lead to enhanced stimulation of the central nervous system and induce euphoria, alertness, reduced appetite, and a sense of increased energy and power. Cardiovascular responses to amphetamine include increased blood pressure and cardiac arrhythmias. More acute responses include anxiety, paranoia, hallucinations, psychotic behavior, and eventually, depression and exhaustion. The effects of amphetamine generally last 2–4 hours, and the drug has a half-life of 9–24 hours in the body. Amphetamine is excreted in the urine in unchanged form and also as hydroxylated and deaminated derivatives.\*

### Principle

The AccuSign\*\*DOA 4 test employs one-step, solid-phase immunoassay technology to discretely detect the presence of any of the above four drugs, or their immunoreactive metabolites, in unine. The assay uses highly specific monoclonal/polyclonal antibodies raised against the target drugs. The test card contains a membrane strip, on which each of the four drugs conjugated to BSA is immobilized at specific locations. The assay is based on the principle of the highly specific immunochemical reactions between antigens and antibodies which are used for the analysis of specific substances in biological fluids. The drug detection relies on the competition for binding to the antibodies between drug conjugates and drugs which may be present in the urine sample.

In the test procedure, a sample of urine is placed in the sample well of the device, and the sample is allowed to migrate upward. If any of the four drugs is present in the urine sample, it forms a complex with the antibody-dye conjugate specific for that drug, and the complex migrates toward the opposite end of the card, passing the specific locations on the membrane where each of the four drug conjugates is immobilized. The drug in the sample competes with the drug conjugate, which is immobilized on the membrane, for the limited antibodies present in the form of antibody-dye conjugate. When a sufficient amount of drug is present, the drug will saturate the antibodies, and the antibody-dye conjugate cannot bind to the drug conjugate on the membrane. Therefore, a drug-positive urine sample will not generate a line at the specific drug position in the result window, indicating a positive result from positive drug competition. Conversely, if a particular drug is absent in the urine specimen, the antibody on the antibody-dye conjugate will bind the membrane-bound drug. In this case, a drug-negative urine sample will generate a line at the specific drug position in the result window, indicating a negative result from an absence of competition with free drug.

Exus.



In addition, the test card has a procedural control built into the system, in the upper control line area. The control line is immobilized with polyclonal anti-mouse antibody; therefore, it will capture monoclonal antibody-dye conjugates that pass the region, showing a colored line in the control (validation) zone. The line works as a procedural control, confirming that proper sample volume was used and the reagent system worked. If insufficient sample volume is used, there may not be a control line, indicating the test is invalid.

### **Materials Provided**

The AccuSign\* DOA 4 test kit contains all the reagents necessary to perform the assay.

- AccuSign" DOA 4 device. The test device contains a membrane coated with drug conjugates in a protein matrix and a pad containing mouse monoclonal anti-THC antibody-dye conjugate, mouse monoclonal anti-opiate antibody-dye conjugate, mouse monoclonal anti-benzoylecgonine antibody-dye conjugate, and polyclonal sheep anti-amphetamine antibody-dye conjugate in a protein matrix.
- Disposable sample dispenser.
- Instructions for use.

### Precautions

- · For in vitro diagnostic use only.
- Avoid cross contamination of urine samples by using a new urine specimen container and dropper for each urine sample,
- This test kit does not contain any HIV or hepatitis infective components. However, urine specimens are potentially intectious. Proper handling and disposal methods should be followed, according to good laboratory practices.
- The AccuSign<sup>1\*</sup> device should remain in its original sealed pouch until ready for use.
- Do not use the test kit after the expiration date.

### Storage and Stability

The AccuSign DOA 4 test kit should be stored at 2-30°C (35-86°F) in the original sealed pouch. The expiration dating was established under these storage conditions.

### **Specimen Collection and Preparation**

Approximately  $150 \,\mu\text{L}$ , of urine sample is required for each test. Fresh urine specimens do not require any special handling or pretreatment. Specimens should be collected in a clean glass or plastic container. If

testing will not be performed immediately, specimens should be refrigerated (2-8°C) or frozen. Specimens should be brought to room temperature before testing.

Specimens containing a large amount of particulate matter may give inconsistent test results. These specimens should be clarified by centrifuging or allowing to settle before testing.

### **Test Procedure**

The test procedure consists of adding the urine sample to the Sample well of the device and watching for the appearance of colored lines in the result window.

### Test Protocol

- For each test, open one AccuSign™ DOA 4
  pouch and label the AccuSign™ device with the
  patient. ID.
- Holding the dropper vertically, dispense 3 full drops (150 μL) of the urine sample into the Sample well.
- 3. Read the result after 5-10 minutes.

### Interpretation of Results

Negative: The appearance of a readish-purple Control line (C) and a line for a specific drug indicates a negative test result: i.e., no drug above the cutoff level has been detected. The color intensities of the Control line and specific drug line may not be equal. A negative test result does not indicate the absence of drug in the sample; it indicates only that the sample does not contain drug above the cutoff level in qualitative terms.

Positive: The appearance of only a reddish-purple Control line and no distinct line next to a specific drug name indicates the test result is positive for that crug (i.e., the specimen contains the drug at a concentration above the cutoff level). A positive lest result does not provide any indication of the level of intoxication or urinary concentration of the drug in the sample; it indicates only that the sample contains drug above the cutoff level in qualitative terms.

Invalid: A distinct Control line (C) should always appear. The test is invalid if no Control line forms at the C position. Such tests should be repeated with a new AccuSign\* DOA 4 test device.

Examples of possible results are shown in the diagram above.

- THC (-), Opiates (-), Cocaine (-), Amphetamines (-): Five reddish-purple lines—one Control line at the C position and one each at the THC, OPL COC, and AMP positions.
- THC (-), Opiates (-), Cocaine (-), Amphetamines (+): Four reddish-purple lines—one Control line at the C position and one line each at the THC. OPI. and COC positions: no line at the AMP position.
- THC (+), Opiates (-), Cocaine (+), Amphetamines (-): Three reddish-purple lines - one Control line at the C position, one line each at the OPI and AMP positions; no lines at the THC and COC positions.
- THC (-), Opiates (+), Cocaine (-), Amphetamines (-): Four reddish-purple lines—one Control line at the C position and one line each at the THC. COC, and AMP positions: no line at the OPI position.
- There are other possible results, depending on the combinations of drugs present in the urine sample.

Note: A very faint line for a specific drug in the result window, visible in 10 minutes, indicates that the amount of drug in the sample is near or below the cutoff level of the test. These urine speciment must be retested, or confirmed with a more specific alternative method such as gas chromatographymass spectrometry, before positive determinations are made.

### Limitations

- The test is designed for use with unadulterated urine only.
- There is a possibility that factors such as technical or procedural errors, as well as other substances in the urine sample than those listed in Table 4 below, may interfere with the test and cause erroneous results.
- Adulterants, such as bleach and/or alum, in urine specimens may
  produce erroneous results regardless of the method of analysis. If
  adulteration is suspected, the test should be repeated with a new
  sample.
- This test detects only the presence of THC metabolites, opiates, cocaine metabolite, amphetamines, and/or their metabolites in urine. A positive test result does not provide any indication of the level of intoxication or urinary concentration.
- The test result read after 10 minutes may not be consistent with the original reading obtained within the 10 minute reading period.
   The test result must be read within 10 minutes of sample application.
- Certain medications containing opiates or opiate derivatives, amphetamines, or methamphetamines may produce a positive result in any chemical or immunological assay. Additionally, foods and tea containing poppy products and/or coca leaves may produce a positive result. Prolonged passive smoking of THC may also produce a positive result.

### User Quality Control

Quality Control: Control standards are not supplied with this kitt however, it is recommended that a control be tested as good laboratory testing practice. NIDA recommends that positive quality control specimens be at or near the cutoff concentration. For information on how to obtain controls, contact PBM's Technical Services. Before using a new kit with patient specimens, positive and negative controls should be tested to confirm the test procedure, and to verify the test produce the expected Q.C. results. Q.C. specimens should also be run anytime there is any question concerning the validity of results obtained.

Process Control: The Control line can be considered an internal process control. A distinct reddish-purple Control line should always appear if the test procedure is performed properly, an adequate sample volume is used, the sample and reagent are wicking on the membrane, and the test reagents are working. If the Control line does not appear

in the control or validation line area, the test is invalid and a new test should be performed. If the problem persists, contact PBM for technical assistance.

### **Expected Values**

AccuSign™ DOA 4 is a qualitative assay. The amount of drugs and metabolites present in urine cannot be estimated by the assay. The assay results distinguish positive from negative samples. Positive results indicate the samples contain the specific drug above the cutoff concentration.

### Performance Characteristics

The AccuSign" DOA 4 Panel Assay detects THC, opiates, cocaine, amphetamines, and their metabolites at cutoff levels based on the recommendations of the National Institute on Drug Abuse (NIDA) for screening of these drugs in urine. 154

THC	11-nor-Δ'-THC-9-carboxylic acid	50 ng/mL
OPI	Morphine	300 ng/mL
COC	Benzoylecgonine	300 ng/mL
AMP	D-Amphetamine	1000 ng/mL

The accuracy of AccuSign" DOA 4 was evaluated in comparison to a commercially available immunoassay (Syva\* EMIT\* II) for each of these four drugs. About 1000 random clinical samples for each drug, including at least 250 positive samples above the cutoff level for each of the four drugs, was tested by both procedures, using the cutoff values listed. Complete agreement was observed in > 99% of the samples as shown below. (Table 1.)

Table 1. Accuracy: Comparison of AccuSign' DOA 4 with Syva\* EMIT\* II Assay

Syva\*EMIT\* II (THC)

		Positive	Negative	TOTAL
AccuSign~	Positive	305	5	310
DOA 4 (THC)	Negative	11	680	691
TOTAL		316	685	1001
	Sy	va* EMIT* I	I (OPI)	
		Positive	Negative	TOTAL
AccuSign"	Positive	249	0	249
DOA 4 (OPI)	Negative	1	716	717
TOTAL		250	716	966
	Sy	va•EMIT• II	(COC)	
		Positive	Negative	TOTAL
AccuSign™	Positive	362	1	363
DOA 4 (COC)	Negative	14	644	658
TOTAL		376	645	1021

Syva*EMIT* II (AMP/MET)			
	Positive	Negative	TOTAL
Positive	185	0	185
Negative	4	291	295
	189	291	480
Relativ	e Sensitivity	Relative S	ecificity
96.59	6 (305/316)		
	Positive Negative Relativ	Positive Positive 185 Negative 4	Positive         Negative           Positive         185         0           Negative         4         291           189         291           Relative Sensitivity         Relative Sensitivity

THC	96.5% (305/316)	99.2% (680/685)
Opiates		
	99.6% (249/250)	> 99% (716/716)
Cocaine	96.3% (362/376)	99.8% (644/645)
Amphetamine	97.8% (185/189)	> 99% (291/291)

In a separate study, AccuSign" DOA 4 was evaluated against specimens confirmed as positive by GCMS, for each of the four drugs. The results below demonstrate the excellent correlation of AccuSign" DOA 4 with GCMS (99% agreement). (Table 2.)

Table 2. Accuracy: Comparison of AccuSign DOA I with GC/MS Assay

		AccuSign™	GC/MS
THC	Positive	87	88
	Negative	1	0
OPI	Positive	73	74
	Negative	1	0
COC	Positive	77	78
	Negative	1	0
AMP	Positive	55	56
	Negative	1	0

### Precision and Accuracy

The precision of the AccuSign" DOA 4 Panel Assay was determined by carrying out the test with serially diluted standard drug solutions. About 98% of the samples containing cocaine, opiates, or amphetamine and about 90% of the samples containing THC concentrations 25% over the cutoff level consistently showed positive results.

The study also included over 40 samples  $\pm$  25% cutoff level as a challenge of cutoff precision. These results were found to be consistently in agreement with expected test results.

### Distribution of Random Error:

Twenty (20) blind samples prepared by spiking various concentrations of cocaine, THC, morphine, or amphetamine were separately tested by two operators. The test results from the two operators showed complete agreement.

### Reproducibility

The reproducibility of the test results of the AccuSign® DOA 4 Panel Assay was examined at three different sites using a total of 15 blind controls, consisting of 5 negative samples, 5 moderately positive samples, and 5 strongly positive samples (i.e., a concentration 3 times the cutoff level). The results obtained at these three sites with these controls demonstrated 100% agreement with each other.

### Specificity

The following table lists compounds that are detected by the AccuSign\* DOA 4 test. The specificity of the AccuSign\* DOA 4 test was determined by adding various drugs and drug metabolities to drug-negative urine specimens and testing with the AccuSign\* DOA 4 test kit. The results are expressed in terms of the concentration required to produce a positive result. (Table 3.)

Table 3. Specificity

Compound	Concentration (ng/mL)	% Cross- reactivity	
THC	( 9)	reactivity	
Cannabinot 11-nor-4'-THC-9-COOH 11-nor-4'-THC-9-COOH -4'-THC -4'-THC OPI	15,000 30 50 25,000 10,000	0,3 160 100 · 0.2 0.5	
Codeine Glucuronide	300 300	100 100	

5 .. -> Hydrocodone 500 60 Hydromorphone 600 50 Levophanol 5,000 6 Meperidine 80,000 0.4 Morphine 300 100 Morphine+3-B-D-glucuronide 500 60 Natorphine 1,000 30 Naloxane 100.000 0.3 Norcodeine 60.000 0.5 Oxycodone 20,000 1.5 Oxymorphone 60,000 0.5 Procaine HCI 100,000 0.3Thebaine 5.000 6 COC Benzoylecgonine 100 Cocaine HCl 500 60 Ecgonine HCl 1,000 30 AMP D-Amphetamine 1,000 100 L-Amphetamine 7,000 14 D.L.Amphetamine sulfate 1,000 100 p-OH-Methamphetamine 30,000 3.3 Methylenedioxyamphetamine 500 200 Methylenedioxymethamphetanine 10.000 10 B-Phenethylamine 20,000 5 Phentermine 5,000 20 Tryptamine 000,001 1 3-OH-Tyramine 90,000 1.1

The following compounds show no cross-reactivity when tested with AccuSign" DOA 4 at a concentration of 100 µg/n/L. (Table 4.)

### Table 4. Non C. oss-Reacting Compounds

Acetaminophen Acetylsalicylaic Aminopyrine Amitriptyline Amotarbital Amoxapine Ampicillin Apomorphine Ascorbic acid Atropine Benzocaine Butabarbital Chlordinazepoxide Chlorpheniramine Chlorpoquine	Dextropropoxyphene Diazepam Diphenylhydantoin Epinephrine Erythromycn Estriol Gentisic acid Glatethimide Guaiacol glycerol ether Hydrochiorothiazude Imapramune Lidocaine Methadone Methadone Methypylon	Naproxen Norethindrone Penicillin Pentobarbital Phencyclidine Phenolbutazone Phenolbutazone Phenolputazone Prednisone Secobarbital Tetracycline Tetrahydrozoline Trifluoperazine Tryptamine Zomepirac

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AccuSign\* is a Trademark of Princeton BioMeditech Corporation.

Patent Pending

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Princeton BioMeditech Corporation P.O. Box 7139, Princeton, New Jersey 08543-7139. U.S.A. 4242 U.S. Route J. Monmouth Junction, New Jersey 08852-1905 U.S.A. Tel: (908) 274–1000 Fax: (908) 274–1010

### AccuSign\*\* BAR

### New One-Step Barbiturates Test

For In Vitro Forensic Use Only

Simple One-Step Immunoassay for the Qualitative Detection of Barbiturates in Urine

### PBM

Catalog No. | DOA-206 | 35 Test Kit | DOA-206 (Q | 16 Test Kit |

### Intended Use

The Accusting BAR test is a simple, one-step, minimumchromatographic assay for the rapid, qualitative detection of barbitarities in arme with a curoff at 300 ng/mL turseconarhatal.

The Accusion BAR (ext provides only a preliminary analysis) result, an are specify: all-maintre extension and be used in order to obtain a confirmat and result. Gas thromatography, man speciosoury (CC(ALS)) is the preferred confirmatory methods are arealable. Chine constitution and preferring doublement should thus result and preferring doublement should be applied on y drag of abuse last result, particularly when preliminary positive traits are used.

### Summary and Principle of Procedure Barburanes are a group of chemicals alcoved from barburanes.

the said Hassified as hypnotics, they deprets the contral nervous system. Taken orally in pill or table form, they are prescribed by many medical conditions, denally for their settative effect. Abuse of bachtimitates con, however, lead in only to impared more coordinations and ment detouch the about respiratory collapses, command feath. The combination of hardwares and also hold in particularly diagnosis. Symptoms of hashinaria abuse include, drowness, shurred speech and irrealistic, Acuse conditions in take expiratory collapse and loss of consecutions. Companions for the chair abuse of consecutions, advances, and death. The effects last at the Acuse Condition of the conditions of the last and Acuse of the Condition of the Condition of the last and acuse of the Condition of the Condition of the last and the Condition of the Condition of the Condition of the last and the Condition of the Condition of the Condition of the last and the Condition of the Condition

### Principle

The Accusign BAR lest uses with phase immunoussus technology for the qualitative detection of secondarhital and barbiturate metabolites in human utine. The test is based on the principle of the highly specific immunichemical reactions between antigens and antihodies which are used for the analysis of specific substance, in histogreal fluids. The test relies on the compension for landing to the ambodies between drift conjugates and dry is which may be present in the nome sample. In the test pro-techne, a sample or name is placed in the sample well of the device of a sample so more is interate upward. If drug is present in the name sample, it competes with the drug compagate, which is munobifized on the membrane, for the limited unabodies present in the form of dye-antibody conjugate. When a sufficient amount of drug or drug metabolite above the cutoff level is present, the drug well saturate the antibodies, thus inhibiting the binding of dye anobady conjugate to the drug conjugate on the membrane. This prevents the formation of a line on the membrane. Therefore, a drug-positive titing simple will not generate a line in the test window, indicating a positive result from positive drug competition, while a negative arms sample will generate a line in the test window, indicating a negative result from an absence of competition with tree drug

In addition to the Teal line that may appear in the Teal window CT, a control line is prevent in the Courant window CT, a control line is prevent in the Courant window CT, or control line viability on the teat. This Control is should always be seen if the easy, conduct out proper came with any appropriate control con

### Materials Provided

The Accusign" BAR test kit contains all the react increasing to perform the assay.

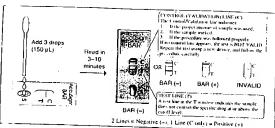
- AccuSign BAR device. The test device contain in membrane control with drug conjugate and apad control golyclond anni-harbiturate ambiody-sive compiliar protein marry.
- Disposable sample dispenser.

### · instructions for use.

### Precautions

- · For in varia forensie use only
- Avoid cross containination of aritie samples by some new more specimen container and dispersion contains sample.
- This test kit does not contain any HIV or h priinfective components. However, near operation.

EXHIBIT 2



- perentially infectious. Proper handling and disposal methods should be followed, according to good laboratory to retices.
- De Accusign<sup>a</sup> device should remain in its original scaled pouch until ready for use.
   De not use the test kit after the expiration date.

### Storage and Stability

AccuSign® BAR test kit should be stored at 2-30°C ("F) in the original scaled pouch. The exputation of the was established under these storage conditions.

### Specimen Collection and Preparation

- (9) symmetry 150 pt. of urine sample is required to such in rech urine speciment do not require any special hair in or porteriament. Specimens should be collected in a compless or plastic consistent. If resting will not be person in immediately, specimens should be chiprecated (2 to 1) or frozen. Specimens should be chrought to ruom temporaries of the consistency specimens and consistency specimens.
- 5 remiens containing a large amount of particulate matter in gryine inconsistent test results. Such specimens should be a sittled by centrifuging or allowing to settle before testing.

### Test Procedure

retest procedure consists of adding the urine sample to the
 myle well of the device and watching for the appearance
 volored lines in the result window.

### Test Protocol

- For each test, open one AccuSign BAR pouch and label the AccuSign device with the patient ID.
- Holding the dropper vertically, dispense 3 drops
   (150 µL) of the urine sample into the Sample
   well (S).
- Read the result after 3 minutes, but within 10 minutes of sample addition.

### Interpretation of Results

Negative: Two Lines The appearance of two reddshippropriate inter-men unter Tex wondow (I) and the once of the Control window (C)—indicates a negative text result to... in barburates above the cutoff level have been descred. The color intensity of the Text lite may be weaker or stronger than that of the Control line. A repative text result does not necessarily window the base level drug in the sample, and indicates the sample does not contain drug above the cutoff feech in qualitative resus.

Pashive: On Line, The appearance of only one reddishparagic line in the Counted window (C) and no district line in the Flast window (T) under the test results presure the the Equivalent C) and the test results presure the the specimen contains bathiusales at a concentration above the cutoff level.) A positive test stand does not provide any indication of the level of invariation or winary concentration of the drug in the sample; it only indicates the sample contains thug above the count flexible in guidative treating

Invalid: A distinct colored (see should always appear in the Control window (C). The test is invalid if no line forms in the Control window (C). Such tests should be repeated with a new Accusing \*\* BAR tests device.

Note. A very faint line in the Test window (T), visible in 10 moutes, indicates that the amount of harmanize in the sample is near or below the casoff level of the test. These sample is near or below the casoff level of the test. These sample is preciment must be received, or confirmed with a more specific alternative method such at gus chromanizy apply must spectrometry, before positive delerminations are must.

### Limitations

- The test is designed for use with unadulterated prine only.
- There is a possibility that factors such as technical or procedural errors, as well as other substances in the urine sample which are not listed in Tables 2 or 3 below, may interfere with the text and cause erroreous results.
- Adulterants, such as bleach and/or alum, in urine specimens may produce erroneous results regardless of the method of analysis. If adulteration is suspected, the test should be repeated with a new sample.
- This test detects only the presence of harbiturates and/or their derivatives in urine. A practice test result does not provide any indication of the level of anoximition or urinary concentration.
- The test result read after 10 minutes may not be consistent with the original reading obtained within the 10 minute reading period. The test must be read within 10 minutes of sample application.
- Certain medicarious containing harbiturates may produce a positive result in any elternical or immunological assay.

### User Quality Control

Quality Control: Control standards are not supplied with that his however, his recommended that a control he trade are well dishrately resting practice. NIDA recommends that a word laboratory resting practice. NIDA recommends that particle quality quarterly specimens be at or near the could concentration. For information in how to obtain controls consider PBMS Technical Services. Before using a new kit withput employee the procedure, positive and negative controls should be leasted to confirm the rest procedure, and to verify the tests probable the aspected QC. results, QC. specimens should also be man unifilm there is any question concerning the validity of results obstanted.

Process Control: The Control line can be considered an interest process control. A distinct reddish-purple Control line should above opposed the text procedure is perhaps upon a first procedure is perhaps upon a distinct process. The sample and reagon are working in the control line does not appear in the control lin

### 

AccuSign\* BAR is a qualitative assay. The amount of secohabital or hiphituate metabolites present in the urine cannot be estimated by the assay. The assay results distinguish positive from negative samples. Positive results indicate the samples contain harbiturates above the cutoff concentration.

### Performance Characteristics

The AccuSign" BAR test has been shown to detect an average of 300 ng/ml of secoharbital in write. The test also detects other barbitals listed below at the minimum concentrations indicated (Table 2)

The accuracy of Accusing a BAR was evaluated in comparison to a commercially available immunoussay (Syva\*EMIT\* D). A unator 302 samples was tested by both procedures. The overall accuracy of the test was 98.75°, as shown below (Table 1.)

Table 1. Accuracy: Comparison of AccuSign\*\* BAR with Sysa\*\* EMIT\*\* 41

### Sysa EMIT H (BAR)

			randarise	TOTAL
AccuSign7	Posture	105	U	105
BAR	Negative	4	193	197
TOTAL		109	193	3/22
AccuSign BAF	Relaine 96 M	Sensitivity (105/109)	Relative Spe 99.9% (193	edieny (193)

### Precision and Accuracy

The precision of AccuSign\* BAR was determined by carrying out the test with senally diluted standard drug solutions. About 98% of the samples containing drug levels 25% over the quintTlevel crossistently showed positive results.

The study also included over 40 samples ± 25% cutoff level as a challenge of cutoff precision. These results were found to be consistently in agreement with expected test results.

### Distribution of Random Error:

Twents (20) blind samples prepared by spiking various concentrations of drug were separately tested by two operators. The test results from the two operators showed complete agreement.

### Reproducibility

The reproducibility of the text neutro of AccoSign\* BAR wave sammed a three dilbreats sick using a total of 15 bind controls. Consisting of 5 neutrons complex, 5 molerately positive samples, and 5 strongly positive samples (i.e., a concentration) threes the conditive level. The results obtained at these these sites with these controls dominate model for 1 strongly and the registering sites at the strongly and t

### Specificity

Compounds that are detected by the Accusing "BAR test are listed below (Table 2). The specificity of Accusing "BAR test are was determined by adding various drugs and this membriles to drug-negative urine specimens and toxing with the Accusing "BAR test kit. The results are expressed in terms of the concentration required to produce a positive result, as

Table 2. Specificity

Compound	Concentration	% Cross-
Allobarbital	200	
Alphenal	1.000	150
Amobarbitat	2,000	30
Aprebarbital		15
Barbitat	200	150
Bulabarbital	2,000	15
	500	60
Butathirat	200	1.50
Cyclopentobarbital	500	
Pentobarbitat	1.000	60
Phenobarbual		30
Secobarbital	5,000	6
. reconstitution	300	100 *

The following compounds show no cross-reactivity when exical with Accussing (BAR attaconcentration of 100  $\mu g/ml$ , (Table § ).

Table J. Non Cross-Reacting Compounds

d-Accessingly and Accessing an	Chlorycon Chlorycon Cholesterel	LEI Egendrine Lei Egendrine Lei Y Eprodine Belle Housepain Bel
--	---------------------------------------	--

Inspirate | Nagrouse | Inspirate | Inspira

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### References

- Hawks RL, Chrang CN, eds. Utime Teiting for Divigit of Abuse Rockwile, MD. National Institute on Drug Abuse (NIDA), Research Monograph 31, 1926.
- Baseli RC, Dispositions of food Drives and Chemicals in Man. Peda Ed., Doors, CA. Bromedt, in Publ. 1982 p. 488.

Accordings" is a Frankristak of Periodenia Bertherhieria Corporations Pale of Pending

Printed in U.S.A.

### PBM

(2) 3.4-Methylicne distaymethamp drawn Methyliphensfate Methyliphensfate Murphine-3-4-Dglucuronale Nalidasie acid Valonghame Nahistone Nahistone Nahistone

Princeton BioMeditech Furparation (F) July 7139 Princeton New Jersey 18543-7139 U.S.A. 123 U.S.Raute, Altonia on Justina, New Jewsy 0852-1985 D.S.A. For 1988 2734-1881

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  Capability of Multi-Drugs

Tests are available in single or multiple test panels for the following drugs:

Amphetamines Barbiturates Benzodiazepines

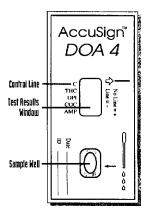
Cocaine Methamphetamines Morphine/Opiates PCP THC (Marijuana)

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EXHIBIT 3

### AccuSign® DOA Series

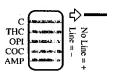
### Test Procedure



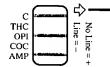
- Using the plastic pipette, add 3 drops of urine sample to the Sample Well.
- 2. Read results in 2 to 5 minutes (within 10 minutes).
- 3. Interpret Test Results Window:
  - CONTROL LIME A colored line indicates
    the test is complete and the system has worked
    properly.
  - NEGATIVE A colored line for the specific drug indicates the test is negative and the drug was NOT DETECTED.
  - POSITIVE No colored line for the specific drug indicates the test is positive and the drug was DETECTED.

Tests Manufactured by Princeton BioMeditech Corporation

### Samples



THC (-) negative Opiates (-) negative Cocaine (-) negative Amphetamines (-) negative



THC (+) positive Opiates (-) negative Cocaine (+) positive Amphetamines (-) negative

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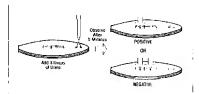
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The tests are composed of preformulated dry reagents arranged on a porous membrane support in a convenient cassette. To perform the tests, just add a few drops of urine to the sample well of the device and wait five minutes. Results are then easily read in the results window as the presence or absence of a red line. A built in reference control ensures that the sample has been added and that the test is effective. The tests are based on the newest, lateral flow micro-particle immunoassay technology and have been evaluated in clinical trials at a major university. The Visualine™II tests are designed to meet NIDA proposed cutoff levels.

<sup>1</sup> A test for Methamphetamine is available for uses that do not require FDA approval to market.



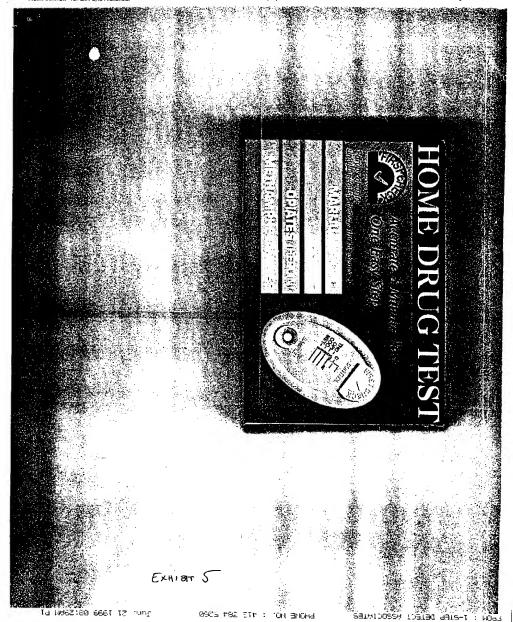
Visualine "Il test device and pipette.

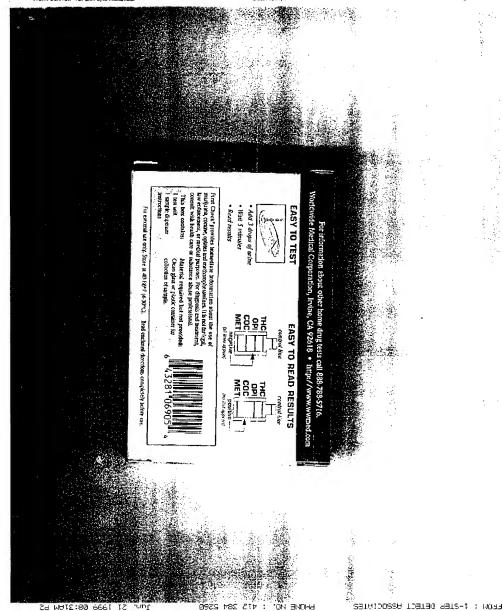


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physician for a recommended liaboratory to relest the same sample Save the unused portion of the unite sample: Ask your family

National Carents' Resource Institute for Drug Education, Inc. (FAIDE Nuri ju ana use increased 43 % in junior high (grades 6-8) and 28% A higher prevalence of depression, medicational problems, and in nork (DAWN), U.S. Dept. of Health and Human Services Sanistational Series: Series I, Number 14-B, Drug Aluse Warning Net caine was mentioned in 55 % and heroitemorphine in 5.3 % of mediin 1994, when manner of drug abuse death was accidental, co was the most commonly reported motive for drug use in 1995. Among energency room cocaine-related episodes, "dependence" Episodes, SAMHSAUMS (August, 1986) 1995 Preliationary Extinutes of Drug-Related Emergency Department 256% between 1991 and 1994 Methaniphelamine-related energeacy department episodes ross sicoppi: University of Mississippi, Resourch Institute of Phurmaceun The potency of marijuans has doubled since the 1970s. Episoden SAMHSAKOAS (August, 1996) department drug related episodes. In 1995, cucaine related episodes comprised 27% of all emergency 9sh annied students' surrey (September 25, 1945) Current Drug Usage Trends Const Franchiston, Marijawa and Today's Youth (1997) terpersonal problems are associated with manipuma use. cal examiner cases. Episodes, SAMHSANAS (August, 1996) 1995 Frelindinary Eminates of Drug-Related Emergency Department "Harrinaver Powercy Monteoring Project Quarterly Report, 1996. Mu 1995 Preliminary Estimates of Drug-Related Energency Departmen Educational Materials ந்தி school students (grades 9-12),

Abust, U.S. Department of Health and Hannan Services, NCATA Pab Isothers His. PHDT12, 1995 Marijuana: Facts Parents Need to Know, National Institute on Drug Department of Health and Human Services, NCADI Publication No Marijuana: Pacts for Teens, benional Institute on Drub Abuse, U.S. Depentury of Health and Human Services, 1995 and Other Caregivers, Conterfor Subspaces Aluse Proceedings, U.S. partition of Education, Bushington, D.C., NCADI Publication No.

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Commonly Asked Questions:

Home Drug Test

Instructions for Use

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control line may look darker than a test line, or vice versa No. The lines need not be the same shade or intensity. The

# They book in a properties and property of the post of the property of the prop

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negative result (5 lines, 1 line in Control window and 4 lines in Test window) will never disappear The test should be read within 10 minutes for best results. A

# land sup diny testresid Mai signal do separation

morphine/opiates, cocaine, and methamphetamines if there The test is working properly as long as a colored line is visible in the Control window. The result is negative for markuana. there is no line next to a drug name in the Test window are also 4 lines in the Test window. The result is positive

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# Do I have to weat the full 5 minutes bottors reading the west 7-53

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Test window), wait 5 minutes but no longer than 10 minutes (1 five in Control window and no line next to a drug name in the test nesults will be clearly reachable in 1 minute. To be sure of a positive your result will be more accurate after 5 minutes. Most negative you do not need to wait. If a line in the test window is not clear lines, a line in the Control window and 4 fines in the lest window) ing the result. However, if your test clearly shows negative (5 Yes, we recommend that you wait the full 5 minutes before read

# What if my child refuses to give a units sample?

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Growing Up Brug Free: A Parent's Guide to Prevention, U.S. De-

Neeping Youth Drug Free: A Guide for Parents, Grandparents,

ty Services listings, or page 4 of this pamphilat for Support under "Alcohol and Drug Abuse" in the Government and Commun sician, your phone directory also provides sources of assistance tinues to refuse, seek professional help. sistant without accusations, threats, or anger. If your child on and why you, as a parent, cannot grove this possibility. Be per phlet supplies current data on drug usage among our youth seriousness of occasional drug usage, page 4 of this pare question the importance of testing him or her for drugs or the that it is their welfare that you are protecting. Should your child Patiently gain your child's cooperation with the understanding Group information and educational resources for more information. Check with your phy-

## Ö Even though the result is negative at still leef that my child night be using drugs. What can't do?

pamphlet for Support Group information. sources of assistance under "Alcohol and Drug Abuse" in the Government and Community Services listings, or page 4 of this Check with your physician, your phone directory also provides

Even through the result is positive, my child claims to be not using drugs. What can I do?

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Methamphetammes Morphine/Opiates, Marijuana (THC), Cocaine, x

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a healthcare or substance abuse professional. medical purposes. For diagnosis and treatment, consult will raetheonpheramines. It is net for legal, law enforcement of the use of marijuana, morphinetopiates, covaine, and Methamphetunines provides immediate information alread First Check<sup>o</sup> Marijuana, Morphine/Opianes, Cacaine, &

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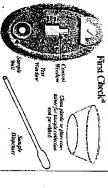
### & Methamphetamines Marijuana (THC), Morphine/Opiates, Cocaine, First Check\* Not to be taken internally.

exposine and remain detectable for 3-10 days after smoking Elevated levels of urinary metabolites are found within thous of in 20-30 minutes and the duration is 90-120 minutes after one cigarette with testawional disorders. The peak effect of smoking manifusors occurfrank toxic delirition. Long term, relatively heavy use may be associated Also, it may cause transient episodes of confusion, anxiety, or even impairment of short term meanwify and manjuana use slows learning When ingested or smoked, it produces explicitly efforts. Users have THC is the printary active ingredient in autojusta (cumabinoids)

## Morphine/Opiates

to normorphiae which is found as a uritary metabolite in both the free (1%) and conjugated (4%) forms. Coderies is excreted as atorphine-6-glucuronide and morphine-3-eilbreal salfate are also is eleminated in the trine as gluculoside. Free morphine in the nrine prototype compound of this group. Up to 15% of the morphine dose pain by depressing the central nerveus system. Morphine is the Opioid analgusies comprise a large group of substances which control an opiale dusc. morphine. Morphine is detectable in the prine for several days after glucuronide conjugate, as free and conjugated narcodeine, and as present. Appoximately 5% of a dose of morphine is N-demethylated accuants for about 19% of the dose, while very small amounts of Heroin is a semisymbelic derivative of marphine.

renzoylergonine in a shun time. Benzoylergonine has a longer bioin a 24 hour period, and excreted in the urine primarity as its more rapid and heightened effect on the abuser. Cocaine is elimi rate, dilusion of the pupits, fever, treuters and swewting. Cornine is nervous system (CNS) stimulant and a local anesthetic Cocoine inurite pH), benznylecgonine (35-54%), and ecgonine (not quantitated) nated in the usine primarity as unchanged drug (1-9%, dependent on "crack" which is especially likely to lead to dependence because of cane base can be smaked in a form that is commonly known as used by smoking, intravenous, intranssal or oral administration. Couser; these psychological effects are accompanied by increased heart duces englicoria, confidence and a sense of increased lenergy in the Comine, derived from the leaves of the cox a plant, is a potent central



gally be detected for 24-60 hours after coming use or exposure. ingical half-life (5-8 hours) than encaine (0.5-1.5 hours) and can gen-

## Methamphetamines

urine for 3-5 days, depending on union pH level. action plactaining use. Methamphetainine is generally detectable in the the presence of the parent compound in the usine indicates However, 10-20% of methamphetamine is exercted unchanged. Thus primarily as ampliciarnine and oxidized and deaminared derivatives! ot 9.24 hears in the body. Methanisherannie is exceeded in the unite methaniphetamine generally last 2-4 hours, and the drag has a half-life and eventually, depression and exhaustion. The effects of responses include anxiety, paragola, halfueinations, psychotic behavior include increased blood pressure and cardiac arrhythmas. More acute energy and power? Cantiovascular responses to methaniphetranile and induce euphroria, alertness, reduced appetite, and a sense of increased higher doses lead to enhanced stimulation of the central nervous system Methamphetamine is a jutent sympathonimselie agent with the agentiapplications. The drug can be taken orally, injected, or inhaled. Acus

## Before you begin

sunlight. Do not use after the expination date stamped on the package low. Store at 36-86' F (2-30' C) in the scaled pouch, away from direc-Read all the information in this pumphlet before per-fountry the research make sure you are familiar with the test kit contents shown, by

### Instructions

Open the seafed pouch a flat surface with Test card, and set the card on da Suloa and Control windows remove the First Check\*



Do not discust the unused write with after the test has been sampleted and the result interpreted & 1875. clean plustic or glassical

With sample disof dispenser tube. sample and release pressure on bulb. Sample will fill half With sample dispenser over sample, press built between thinih and index higer, liver dispenser opening into



Allow the test card to remain undisturbed until result is read. Read the result after 5 minutes but within 10 minutes -

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### Positive

dow means the sample contains that drug Out line in the Control window and no line next to a drug name in the Test win





appear in the Control (upper) window The test instructions must be followed precisely. Limitations do not interpari result If no line appears in the Centrol window

The First Checks One-Step Home Drug Test is not reusable

### lites in urine. A positive test dues not provide any internation phine/opiates, cocaine, and methamphetumines or their metahoabout the autount or level of intoxication The test detects only the presence of manifecta (TEC), then

should be repeated with a new urine sample duce an erroneous result. If adulteration is suspected the sea terants, such as bleach and/or alum, in a urine sample may pro The test is designed for use with imadulterated mine, only. Actual The result must be read 5-10 minutes after sample application. A

Urine sample should be at room temperature. If sample has been result read after 10 minutes may not be accurate.

refrigurated, allow sample to reach room temperature before testing

ordhand manijuma smoke may produce a posupre result Certain medications containing opiates or methamphetamines may leaves may jweeduce a positive result. Frolonged expersum to see Additionally, floods and tea containing poppy products and/or coca produce a positive result in any chemical and in ununological assay

### Kesults

Wait at least 5 minutes but not more than 10 minutes before

reading result



ny ou se ates, eocaine, nor methamplietamine there is no marijuona, morphine/cpi

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no drug taken

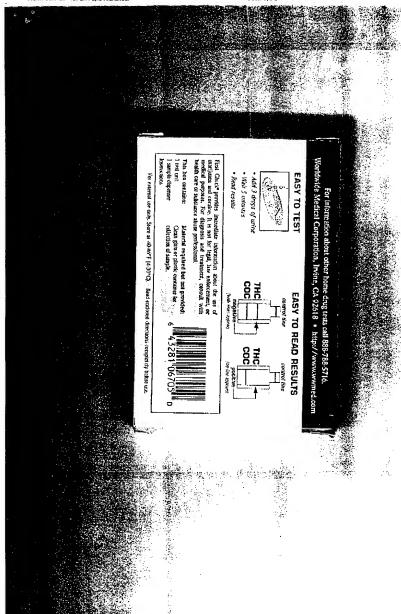
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PAGE 25/33 \* ROVD AT 9/26/2004 2:45:43 PM (Eastern Daylight Time) \* \$VR:USPTO-EPXRF-3/26 \* DNIS;2/31254 \* CSID:Dan Sullivan Jr. \* DURATION (mm-ss):22-32



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Page 27 of 33



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Marijaan Paency Mondoring Project: Quarterly Report: 1906, Mis The potency of marijuana has doubled since the 1970s Episodus SAMHSARONS (August, 1996) 1995 Preliminary Estimates of Drug-Related Emergency Department department drug related episodes. in 1995, country related episodes comprised 27% of all emergency

attention, a decreased capacity to shift attention, reduced learning Pape H. and Yangelun-Toold D. Journal of the American Medical Asso. and decreased mental flexibility. Heavy marijuans use is associated with reduced ability to sastain sissippi: University of Mississippi, Research Institute of Pharmacout Call N. P. W. C.

Episodes: SAMHSANDAS (August, 1984) Among emergency room creatur-related episodes, "dependence cimios. 275, 521-527 (1996) was the most commonly reported modifie for drug use in 1995. 1995 Preferency Edinades of Drug Related Emergency Department

work (DAWN), U.S. Dept. of Health and Hunton Services Sunistivital Series: Series L. Number 14-B. Drug Abuse Warning Net caine was mentioned in 55% of medical examiner coses. In 1994, when manner of drug abuse death was accidental, on

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Caron Foundainn, Marijuwas and Today's Youth (1997) terpersonal problems are associated with marijuuns use. A higher prevalence of depression, motivational problems, and

permant of Education, Withington, D.C., NCADI Publication Growing Up Drug Free: A Parent's Guide to Prevention, U.S. Educational Materials

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Department of Hruthi and Human Services, 1995 Keeping Youth Drug Free: A Guide for Parents, Grandparents and Other Caregivers, Center for Substance Alasse Provintion, U.S.

Department of Health and Hunart Services, NCAM Publication No. Marijuana: Facts for Terns, National Institute on Dub Abust. U.S.

Marijuara: Facts Parents Need to Know, National Institute on Dray Abase, U.S. Department of Health and Human Services, NCAHI Fub hration No. FHD712, 1995

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# Commonly Asked Questions

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- No. The lines need not be the same shade or intensity. The control line may look darker than a test line, or vide versa

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# Hoping on the optical particular and an experience of

negative result (3 lines, 1 line in Control window and 2 lines in Test window) will never disappear. The lest should be read within 10 minutes for best results.

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is positive if there is no line next to a drug mame in the Test cocaine if there are also 2 lines in the Test window. The result in the Control window. The result is negative for manuage and The test is working properly as long as a colured line is visible

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(1 Fre in Control window and no line next to a drug name in the test results will be dearly readable in 1 minute. To be sure of a positive you do not need to wait. If a line in the fest window is not clear lines, a line in the Control window and 2 lines in the test window) ing the result. However, if your test clearly shows negative (3) Yes, we recommend that you wait the full 5 minutes before read Test window), wait 5 minutes but no longer than 10 minutes your result will be more accurate after 5 minutes. Most regalive

# Main the contraction of the contract of the co

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ty Services listings, or page 4 of this paraphlet for Support Group information and educational resources for more information. Be persistent without accusations, threats, or enger. If your child of this pamphlet supplies current data on drug usage among our youth and why you, as a parent, cannot ignore this possibility. question the importance of testing him or her for marijuana and that it is their welfare that you are protecting. Should your child Patiently gain your child's cooperation with the understanding continues to reluse, seek professional help. Check with your accaine or the seriousness of occasional drug usage, page 4 under "Alcohol and Drug Abuse" in the Government and Commun physician, your phone directory also provides sources of assistance

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Check with your physician, your phone directory also provides pamphlet for Support Group intermetion Covernment and Community Services tistings, or page 4 of this sources of assistance under "Alcohol and Drug Abuse" in the

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physician for a reconstrended liaboratory to reflect the same sample Save the unused portion of the urine sample. Ask you lamily

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## HOME DRUG

Instructions for Use



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diagnosis and treatment, consult with a health care o not for legal, law enforcement, or medical purposes. Fin information about the use of marifuana and covaine. First Checks Marijuana & Cocaine provides immediate substance abuse professional.

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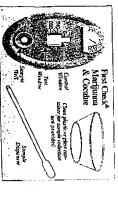
## Viol to be taken internally First Checke Marijuana & Cocaine



# Marijuana & Cocaine

memory and manjuana use slows learning. Also, it may euphoric effects. Users have impairment of short term cause transient episodes of confusion, anxiety, or even (cannabinoids). When ingested or smoked, it produces boars of exposure and remain detectable for 3-10 days Elevated levels of urinary metabolites are found within and the duration is 90-120 minutes after one eigerette effect of smoking marijuana occurs in 20-30 minutes mny be associated with behavioral disorders. The peak frank toxic delirium. Long term, relatively heavy use THC is the primary active ingredient in marijuana

oral administration and excreted in the urine primarily Cocaine is used by smoking, intravenous, intranasal or rate, dilation of the pupils, fever, tremors and sweating. chological effects are accompanied by increased heart and a sense of increased energy in the user; these psypotent central acryous system (CNS) stimulant and a local anesthetic. Coraine induces euphoria, contidence as benzoyleegonine in a short time. Benzoyleegonine Comine derived from the leaves of cora plant is a



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many after cocaine use or exposure (0.5-1.5 hours) and can generally be detected for 24-60 day is marger monageness that the the transcription of a transcription is a transcription of the transcription of

## Before you begin

the test kit contents shown, below. Store at 36-86' F (2) not use after the expiration date stamped on the package. (i) C) in the sealed pouch, away from direct sunlight. Do forming the test. First, make sure you are familiar with Read all the information in this partiphlet before per

## Instructions

 Open the sealed pouch, remove the First Check\* card, and set the card on a flat surface with Test and Control wardows lucing up.

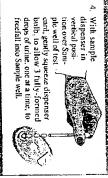
Positive

Sino drug taken

- Collect urine sample in a clean plastic or glass container
- With sumple dispenser over sample. press bulb between

س:



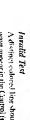


Allow the test eard to remain undisturbed unutes but within 10 minutes til result is read. Read the result after 5 min-









window. If no line appears in the Control window, do not ways appear in the Control (upper) A distinct colored line should alinterpret result





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## Limitations

usuble. The test instructions must be followed precisely The First Check\* One-Step Home Drug Test is not re

and cocaine or their metabolites in urine. A positive less does not provide any information about the amount or The test detects only the presence of marijuana (THC) level of infoxication

tion. A result read after 10 minutes may not be accurate. pected, the test should be repeated with a new trine sample may produce an erroneous result. If adulteration is sus Adulterants, such as bleach and/or aham, in a utine sampl The test is designed for use with unadulterated urine, only The result must be read 5-10 minutes after sample applica

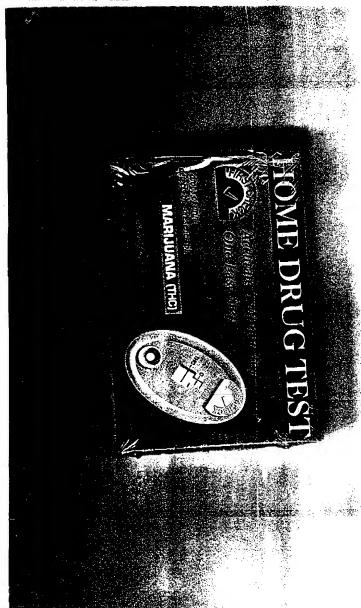
perature before testing. has been refrigerated, allow sample to come to rown tem-Unine sample should be at room temperature. If sample

produce a positive result Prolonged exposure to secondhand marijuana smoke may

### Kesnus

i glare reading sesuit. Wait at least 5 rainates but not priore than 10 minutes

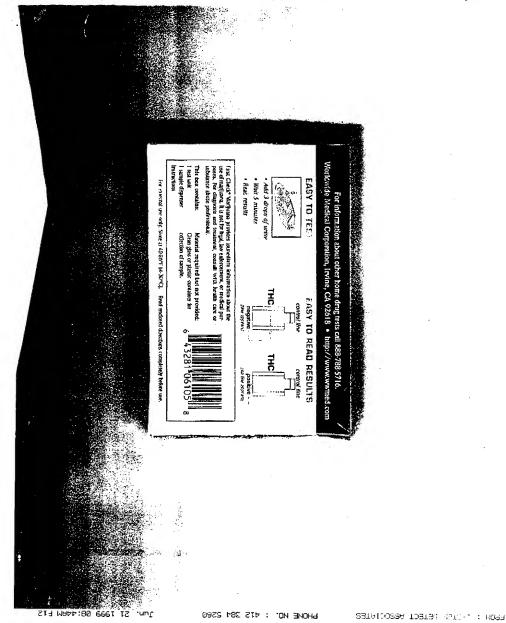




n. 21 1999 08:42AM P11

ME NO: : 415 384 2560

FROM : 1-STEP DETECT ASSOCIATES



Culver City, CA 90231-3475 800/736-9805 Families Anonymous, Inc. PO. Box 3475 5. annual students' survey (September 25, 1996) lationed Parents' Resource Institute for Drug Education, Inc. (I'RIDE) high school students (grades 9-12). lorijuana use increased 43% in Junior bigh (grades 6.8) and 28% urrent Marijuana Tremts

attention, a decreased capacity to shift attention, reduced harming and decreased mental flexibility. type H. and Bergeton. Fuld D., Journal of the American Medical Astrayy marijuana use is associated with reduced ability to sustain issippi University of Mississuppi, Research Institute of Pharmaceuti-Marijuana Patency Musikering Project: Quarterly Report, 1996, Mis the potency of marijuana has doubled since the 1976s

(bap these who never use marijuana. at least 3 times in the last month, are less likely to use condons metation, 275, 321-527 (199a) Ope marijuans eigarette equals (wenty regular eigarettes in terms New Rod: Columbia University. Center on Addition and Substance 15% and 25% of boys and girls, respectively, who used marijauno Jashbin D.P. and et al. National Institute on Drigs and Alcohol Red brokemal damage. Abuse: [1996, Jose,

starch Managraph, 99 (1990) Canon Foundation, Marijuana and Today's Youth (1997) terpersonal problems are associated with marijuana use. A higher prevalence of depression, multivational problems, and  $\hat{m}$ Educational Materials

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Keeping Youth Drug Free: A Guide for Parents, Grandparents Growing Up Drug Free: A Parent's Guide to Prevention, U.S. parented of Edworton, Washington, D.C., NCAIN Publication Department of Health and Haman Services, NCADI Publication 6 Marijuana: Facts for Teers, National limitate on Diph Almee, U.S. Department of Health and Historia Services, 1995 and Other Caregivers, Center for Substance After Prevention, U.S. 3

lication No. PHD712, 1995 Abuse, U.S. Department of Health and Human Services, NCADI Pa Marijuana: Facts Parents Need to Know. National Institute on Dr.

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10013-0862

P.O. Box 862

Al-Anon Family Group Headquarters

Support Groups

PO. Box 2012 Marijnana Amanymous World Services Office

1001/766-6779 Van Nuys, CA 91404

÷ physician for a recommended laboratory to refest the same Save the unused portion of the unine sample. Ask your family

Visit Worldwide Medical Corporation at http://www.ewwgned.com

MOSTDINIDE WEDICY Living CA 92618 U.S.A soft free 888/788.5716

# Commonly Asked Questions:

HOME DRUG'HEST

Instructions for Use

- mean some marijuana is present? The test time is lighter than the line in the Control window. Does this
- control line may look darker than the test line, or vice virsa (to. Buth lines need not be the same shade or intensity
- How long will the colored line remain visible?

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- twe result (2 lines, 1 line in Control window and 1 line in Test windown will never disappear The test should be read within 10 minutes for best result. A mega
- lam not sufe of my test result. What should I do?

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- The test is working properly as long as a colored line is welche in the Control window. The result is negative for marijuena if there is also a line in the Test window. The result is positive in there is no line in the Test window.
- Do I have to want the fall 3 minutes belove reading the lest?

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- sult will be more accurate after 3 minutes. Most nagative test ines, a line in both the control and test windows), you do not ing the result. However, if your test clearly struis negative(2 Yes, we recommend that you want the full 3 minutes before read but no longer than 10 minutes. line in Control window and no line in Test window), wit 3 minutes results will be clearly readable in 1 minute. To be sure of a positive (1 need to wait. If the line in the test window is not clear, your re-
- What if my child refuses to give a urine sample?

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- Patiently gain your child's cooperation with the understanding Group Information and educational resources for more information munity Services listings, or page 4 of this pamphic for Suppor under 'Alcohol and Drug Abuse' in the Government and Comcian, your phone directory also provides sources of assistance signes to refuse, seek professional help. Check with your physipersistent without accusations, threats, or anger. If your child comyouth and why you, as a parent, cannot ignore this possibility. pamphiet supplies current data on merijaana usaga among out the senousness of occasional marijuana usage, page 4 of this question the importance of testing him or her for manifusna or that it is their wettare that you are protecting. Should your child æ
- Ever though the receiling had allowed a sill beet that my child then be using door. When control the first and the state of the sill be seen to pamphlet for Support Group information Government and Community Services listings, or lage 4 of this Check with your physician, your phone directory also provides sources of assistance under "Akonhol and Drug Ahuse" in the

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### Marijuana First Check® (THC)

When the need to know is....now

Simple - one step

- Easy-to-reac
- Confidentia
- Result in 3 minutes

enforcement, or medical purposes, for discovering about the use of morijuana. It is not be lexish her professional treatment, counds with a health care or substices conv First Check Marijuana provides immediate extremation

## Read the following directions completely before use.

Store at 36-86°F (2-30°C) For external use only.

# FOR EDUCATIONAL USE ONLY

First Check" is a metatered trusteened of Worldard: Medical Company

Not to be taken internally

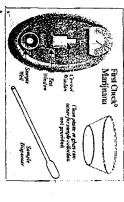


## Marijuana (THC)

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## Before you begin

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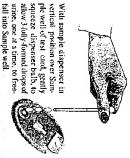
## Instructions

- 1. Open the sealed pouch, remove the First Check! and Control windows facing up. card, and set the card on a flat surface with Text
- Collect urice sample in a clean plastic or glass container.



With sample dispenser over sumsure on bulb. You should see saming into sample and release presple, press bulb between thumb and pie fill half of dispenser tube. index finger insert dispenser open-

fest has been completed and the result Do not discard the united urine until



Allow the test card to remain undisturbed until result is read. Read the result after 3 minutes but within 10 minutes

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# to not use after the expiration date stamped on the pack

# 2-30 C) in the scaled pouch, away from direct sunlight



I line - positive



sample contains manjuana

drug taken A distinct colored line should al-



### Invalid Tes

window. If no line appears in the Control window, do not ways appear in the Control (upper)

## Limitations

interpret result

usable. The test instructions must be followed precisely The First Check" One-Step Home Drug Test is not in

window) does not provide any information about the its metabolites in urine. A positive test (no line in Test The test detects only the presence of manipuana (THC) or amount or level of intoxication.

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produce a positive result Prolonged exposure to secondhand marijuana smoke may

### Results

before reading result What at least 3 minutes but a cone than 10 amoutes



dow may be lighter or durker than sample. The line in the Test win is no marijuana present in the urine Test (lower) window, means there of the Control (upper) window are Two horizontal lines, one in each 2 lines - negative